

CHIRAL CHEATING AGENT AND CHIRAL CATLYST

CROSS-REFERENCE TO RELATED APPLICATION

5 This application claims the priority benefit of Taiwan application serial no. 92104138, filed February 27, 2003.

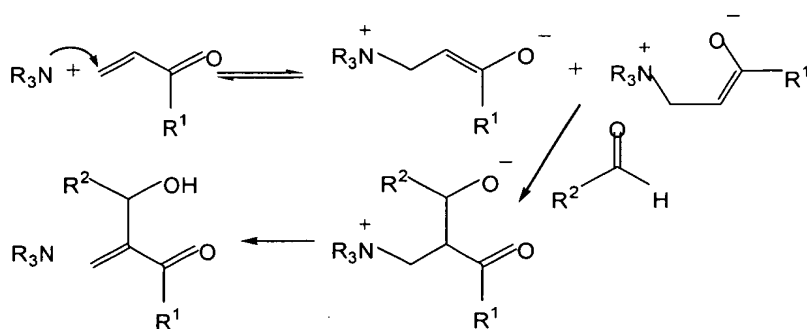
BACKGROUND OF THE INVENTION

Field of the Invention

10 **[0001]** The present invention relates to chiral cheating agents and chiral catlysts, wherein the chiral catlysts are formed from the chiral chelating agents and metals.

Background of the Invention

[0002] The use of catalytic asymmetric reactions for the synthesis of enantiomerically pure compounds has been an ongoing effort of organic chemists. The coupling of an α,β -unsaturated carbonyl/nitrile with an aldehyde (Morita-Baylis-Hillman reaction) produces a useful functionalized acrylate that allows for further functional group manipulation. The reaction can be catalyzed by using a chiral catalyst to synthesize a product having high level of enantiomeric excess. The mechanism of the Baylis-Hillman reaction, which is mediated by a tertiary amine, is illustrated as
20 follows:



Although the diastereoselective Baylis-Hillman reaction has been shown to proceed, in some cases, with high to excellent diastereoselectivities, the enantioselective variation of this reaction is less well developed.

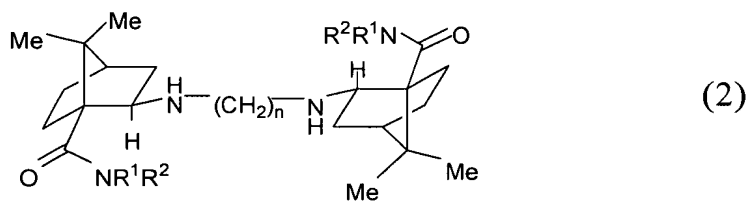
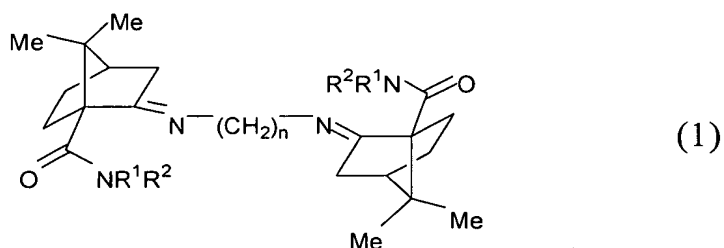
- 5 **[0003]** Recently, Hatakeyama et al. reported that a high to excellent enantiomeric excess can be achieved when quinidine derivatives are used as chiral amine catalysts (*J. Am. Chem. Soc.*, **1999**, 121, 10219). The use of a chiral Lewis acid to catalyze this transformation is a common strategy. Aggarwal and co-workers report on the use of lanthanides and group III metal triflates to accelerate the Baylis-Hillman
- 10 reaction (*J. Org. Chem.* **1998**, 63, 7183 and *J. Chem. Soc. Chem. Commun.*, **1996**, 2713). However, only 5% ee was obtained, when these metals are complexed with a broad range of oxygen-rich chiral ligands. No practical levels of enantioselectivity have yet been reported when a chiral Lewis acid catalyst is used.

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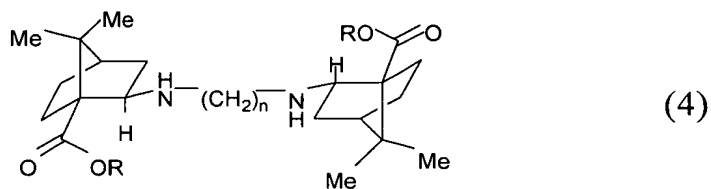
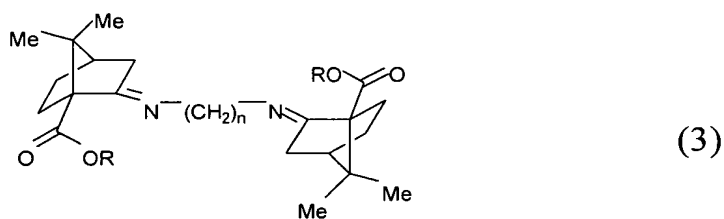
SUMMARY OF THE INVENTION

[0004] The present invention provides chiral chelating agents and chiral catalysts, which are formed from the chiral reagents and metals to improve the enantioselectivity of catalytic asymmetric reactions.

[0005] The present invention provides a chiral chelating agent, which is a camphor derivative having a structure of any one of the formulas (1) to (17) and an enantiomeric or an diastereomeric isomer of any one of the formulas (1) to (17):

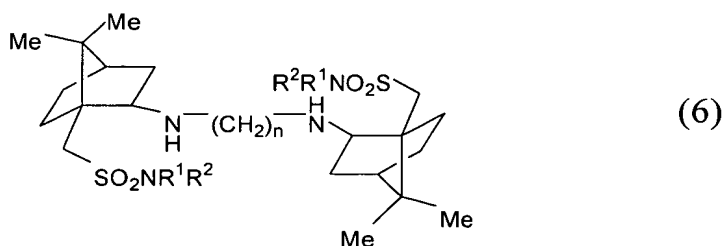
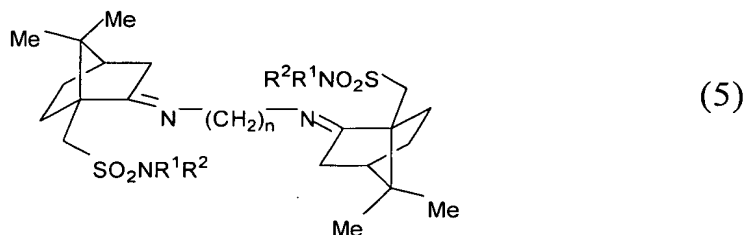


wherein R¹ and R² represent H, methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic group, an aromatic group substituted at 2-, 3- or 4-position, an aromatic-like group, or a naphthyl or naphthyl-derived group, and n is an integer between 0 and 4;

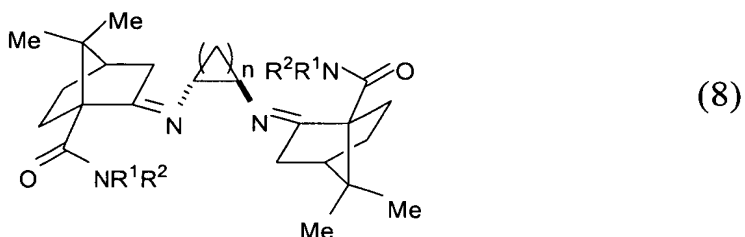
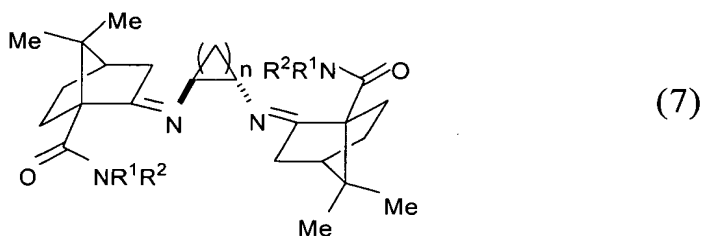


wherein R represents H, methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic

group, an aromatic group substituted at 2-, 3- or 4-position, an aromatic-like group, or a naphthyl or naphthyl-derived group, and n is an integer between 0 and 4;



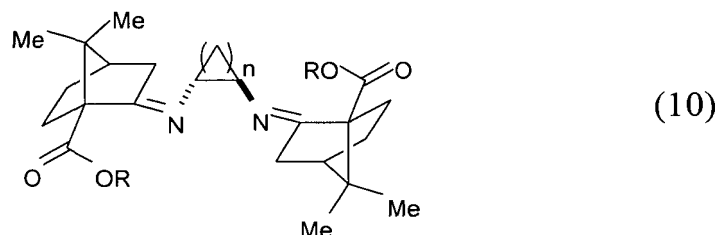
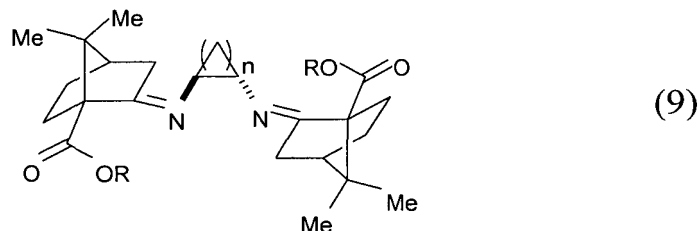
5 wherein R^1 and R^2 represent H, methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic group, an aromatic group substituted at 2-, 3- or 4-position, an aromatic-like group, or a naphthyl or naphthyl-derived group, and n is an integer between 0 and 4.



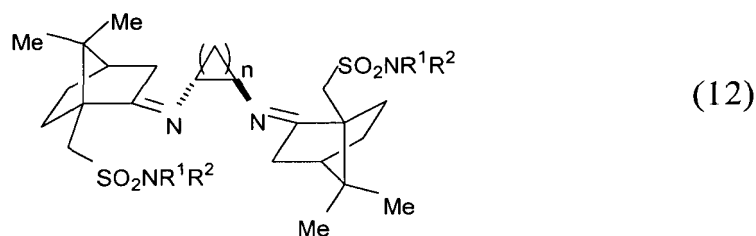
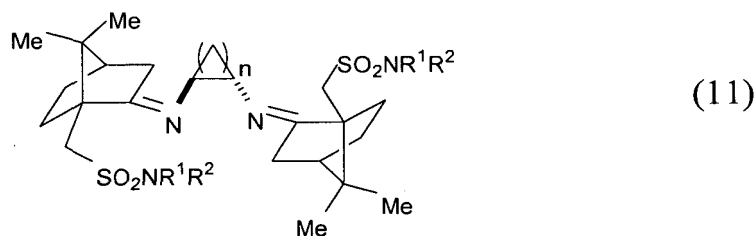
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wherein R^1 and R^2 represent H, methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic

group, an aromatic group substituted at 2-, 3- or 4-position, an aromatic-like group, or a naphthyl or naphthyl-derived group, and n is an integer between 0 and 4.



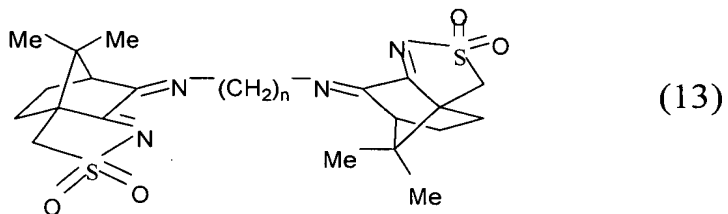
- 5 wherein R represents H , methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic group, an aromatic group substituted at 2-, 3- or 4-position, an aromatic-like group, or a naphthyl or naphthyl-derived group, and n is an integer between 0 and 4.



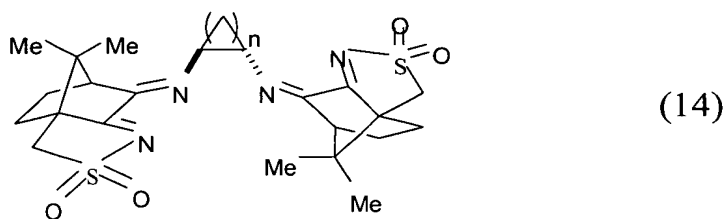
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wherein R^1 and R^2 represent H , methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic

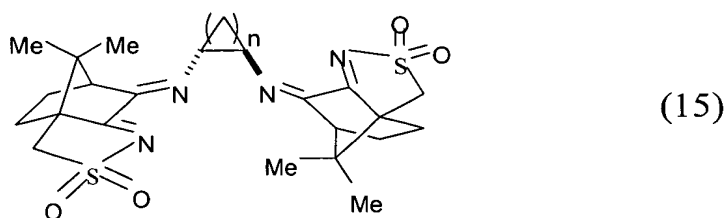
group, an aromatic group substituted at 2-, 3- or 4-position, an aromatic-like group, or a naphthyl or naphthyl-derived group, and n is an integer between 0 and 4.



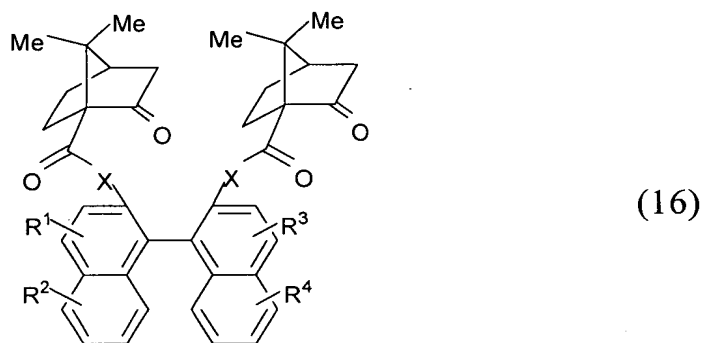
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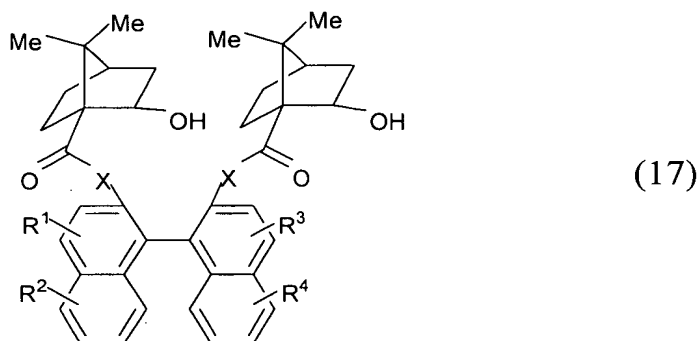


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wherein n is an integer between 0 and 4.





wherein X represents an oxygen atom or a nitrogen atom; R¹, R², R³ and R⁴ represent H, methyl, ethyl, a primary, secondary or tertiary straight, branched or cyclic alkyl group having 3-7 carbon atoms, a heterocyclic or aromatic group, an aromatic group substituted at 2-, 3- or 4-position, a naphthyl or naphthyl-derived group or the above groups substituted with at least one halogen.

[0006] In addition, the chiral chelating agents of this invention comprise the diastereomers of the formulas (16) and (17).

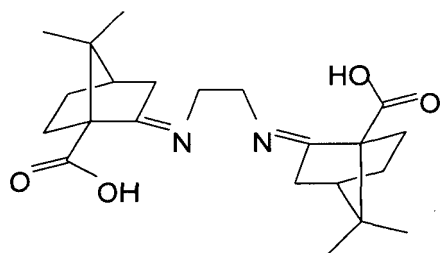
[0007] The present invention provides a chiral catalyst, which is formed from the chiral chelating agent of any one of the above formulas (1) to (17) and a metal, wherein the metal comprises an alkali metal, an alkaline earth metal or a transition metal including a lanthanide metal.

[0008] The chiral catalyst of this invention is suitable for use in a Baylis-Hillman reaction. In some cases, the enantiomeric excess of the product is determined to be 71%. This represents that the chiral catalysts of this invention improve the enantioselectivity of catalytic asymmetric reactions.

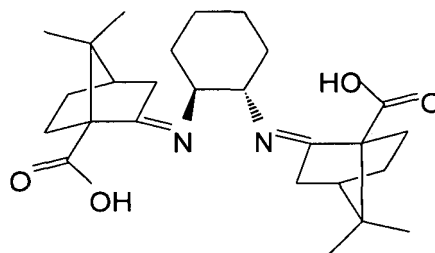
[0009] It is to be understood that both the foregoing general description and the following detailed description are exemplary, and are intended to provide further explanation of the invention as claimed.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

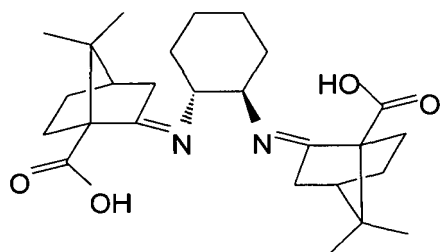
[0010] The chiral chelating agents used in this embodiment comprise four agents, and the chiral castlysts used in this embodiment are formed from the four agents and lanthanide metals. The four agents is selected from the chiral chelating agents as illustrated above in formulas (3), (9) and (10), wherein the structures of the chiral chelating agent 1 and 4 are the above formulas (3) when R is H, and n is 2 and 0, respectively. The structures of the chiral chelating agent 2 and 3 are the above formulas (9) and (10) when R is H, and n is 4, respectively.



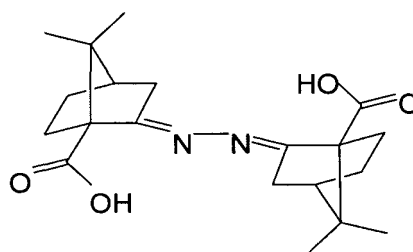
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The four chiral chelating agents are prepared from the condensation of (+)-ketopinic acid with the corresponding diamines or hydrazine under acidic conditions. The chiral

chelating agents 1 and 4 can be easily synthesized using ethylenediamine and hydrazine in the presence of acetic acid, and the yields are 69% and 88%, respectively. The chiral chelating agents 2 and 3 are provided by treatment of racemic *trans*-1,2-diaminocyclohexane with (+)-ketopinic acid in refluxing CHCl_3 for 36 hours. That the
 5 chiral chelating agents 2 and 3, which are diastereomers, can be separated by flash column chromatography and the absolute stereochemistry of the chiral chelating agents 3 is further confirmed by single crystal X-ray analysis.

[0011] The chiral chelating agent 1 is produced as follows: first, ethylenediamine (1.64 g, 27.4 mmol) and acetic acid (0.1 mL) is added into a solution
 10 of (+)-ketopinic acid (10 g, 54.9 mmol) in CHCl_3 (100 mL) at room temperature. The resulting mixture is then refluxed for 36 hours and the reaction then quenched with H_2O (50 mL). The resulting solution is extracted with CH_2Cl_2 (100 mL) and the layers are separated. The organic layer is washed with brine (10 mL), dried (MgSO_4) and concentrated. The crude product is purified by silica gel using $\text{EtOAc}/\text{CH}_2\text{Cl}_2$ as the
 15 effluent (4/1) to give 7.22 g (69%) of the chiral chelating agent 1 as a white solid. The product is analyzed and the data is as follows:

^1H NMR (CDCl_3 , 200 MHz) δ 3.65 (dd, $J = 15.0, 3.8$ Hz, 4H), 2.56 (dd, $J = 18.2, 3.6$ Hz, 2H), 2.40 (td, $J = 12.2, 3.6$ Hz, 2H), 2.15-1.97 (m, 6H), 1.68 (td, $J = 12.8, 4.0$ Hz, 2H), 1.43-1.30 (m, 2H), 1.25 (s, 6H), 0.87 (s, 6H); ^{13}C NMR (CDCl_3 , 50 MHz) δ 185.09, 173.21, 60.68, 50.78, 50.30, 43.86, 35.40, 31.62, 27.84, 20.07, 19.85; MS m/z (relative intensity) 388 (M^+ , 8), 373 (10), 319 (100), 163 (23), 148 (30); HRMS m/z 388.2336 (calcd for $\text{C}_{22}\text{H}_{32}\text{N}_2\text{O}_4$ 388.2362).

[0012] The chiral chelating agent 2 and 3 are produced approximately the same way as above-mentioned except racemic *trans*-1,2-diaminocyclohexane having an

optimum amount is in place of ethylenediamine. The products are analyzed and the data is as follows:

The chiral chelating agent 2:

¹H NMR (CDCl₃, 200 MHz) δ 3.45 (m, 2H), 2.52 (m, 2H), 2.36 (m, 2H), 2.11-1.98 (m, 6H), 1.81-1.67 (m, 4H), 1.60-1.24 (m, 6H), 1.22 (s, 6H), 0.86(s, 6H); ¹³C NMR (CDCl₃, 50 MHz) δ 183.34, 173.13, 64.38, 60.19, 50.37, 43.68, 34.99, 31.77, 30.54, 27.84, 23.76, 20.04, 19.57; and HRMS m/z 442.2834 (calcd for C₂₂H₃₂N₂O₄ 388.2362).

The chiral chelating agent 3:

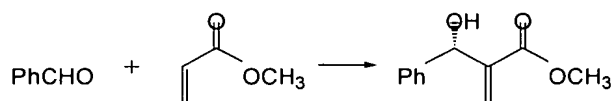
¹H NMR (CDCl₃, 200 MHz) δ 3.43 (dd, J = 5.2, 3.2 Hz, 2H), 2.61 (t, J = 3.0 Hz, 1H), 2.54 (t, J = 3.8 Hz, 1H), 2.35 (td, J = 12.1, 4.6 Hz, 2H), 2.11-1.92 (m, 6H), 1.83 (d, J = 6.6 Hz, 2H), 1.65-1.27 (m, 10H), 1.25-1.24 (m, 2H), 1.23 (s, 6H), 0.83 (s, 6H); ¹³C NMR (CDCl₃, 50 MHz) δ 183.24, 173.10, 64.33, 60.55, 49.31, 43.93, 34.93, 32.41, 31.07, 27.90, 23.85, 20.17, 20.06; MS m/z 442 (M⁺, 7), 262 (35), 217 (50), 148 (45), 81 (100); HRMS m/z : 442.2833 (calcd for C₂₂H₃₂N₂O₄ 388.2362); and Crystal data at 22°C: C₂₆H₃₈N₂O₄ M 442.59, tetragonal, P4₃2₁2, a = 7.8028(13) Å, c = 40.282(4) Å, V = 2452.5 Å³, Z = 4, λ = 0.70930 Å, F(000) = 960.56, D_c = 1.199 Mg/m³, μ = 0.08 mm⁻¹, 5242 reflections, 146 parameters, R = 0.051, and R_w = 0.067 for all data.

[0013] The chiral chelating agent 4 is produced approximately the same way as above-mentioned except hydrazine is in place of ethylenediamine and CH₂Cl₂ is in place of CHCl₃. The products are analyzed and the data is as follows:

¹H NMR (CDCl₃, 200 MHz) δ 2.86 (t, J = 3.6 Hz, 1H), 2.74 (t, J = 3.8 Hz, 1H), 2.53 (td, J = 12.6, 4.8 Hz, 2H), 2.32-2.04 (m, 6H), 1.82 (td, J = 9.0, 4.2 Hz, 2H), 1.45 (m, 2H), 1.29 (s, 6H), 0.98 (s, 6H); ¹³C NMR (CDCl₃, 50 MHz) δ 181.90, 171.84, 60.99, 51.74, 44.03, 35.63, 31.65, 27.69, 20.01, 19.89; MS m/z 360 (M⁺, 10), 316 (40), 288 (60), 163

(75), 148 (100), 134 (80), 95 (90); HRMS m/z 360.2011 (calcd for $C_{22}H_{32}N_2O_4$ 388.2362) Crystal data at 22 °C: $C_{20}H_{28}N_2O_4$ M 360.45, monoclinic, C2, $a = 11.414(3)$ Å, $b = 7.580(4)$ Å, $c = 21.887(3)$ Å, $V = 1865.4$ Å³, $Z = 4$, $\lambda = 0.70930$ Å, $F(000) = 776.48$, $D_c = 1.283$ Mg/m³, $\mu = 0.09$ mm⁻¹, 1812 reflections, 235 parameters, $R = 0.042$, and $R_w = 0.079$ for all data.

[0014] The lanthanide metal triflates were screened with the synthesized chiral chelating agents 1-4 using benzaldehyde and methyl acrylate (A), i.e. Baylis-Hillman reaction as model probes.



The catalytic system used in this case has a number of appealing features, which include (i) the molar ratio of chiral ligand to metal salts is designated to be 2:1 so that optimum enantioselectivity could be obtained under such conditions, deviation in either direction led to a decrease in selectivity (data not shown); (ii) a catalytic amount of catalyst (3 mol% of metal) is sufficient for asymmetric induction; (iii) the complexes are completely soluble in CH_3CN , giving a homogeneous solution in that ratio; and (iv) to prevent the formation of potential amine-Lewis acid complexes and therefore weaken the nucleophilic ability, a strictly controlled amount of 1,4-diazabicyclo[2.2.2] octane (DABCO) (10 equiv to the catalyst or 30 mol% to substrate) is used. The result is shown in Table 1 as follows:

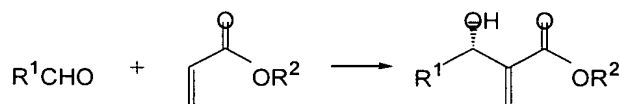
Table 1 Reaction of benzaldehyde and methyl acrylate ^a

entry	ligand	Lewis acid	yield(%) ^b	%ee ^c	configuration ^d
1	1	Eu(OTf) ₃	81	0	S
2	1	Yb(OTf) ₃	72	17	S
3	1	La(OTf) ₃	75	84	S
4	2	La(OTf) ₃	70	67	S
5	3	La(OTf) ₃	71	11	S
6	4	La(OTf) ₃	0	-	-

^a Benzaldehyde (0.5 mmol) is reacted with methyl acrylate A (0.5 mmol) in the presence of DABCO (30 mol%), Lewis acid catalyst (3 mol%) in CH₃CN (2.6 mL) at room temperature for 10 hours. ^b Isolated yield, ^c Determined by HPLC analysis using a chiral column, ^d Determined by comparison of optical rotation with the literature value.

[0015] From Table 1, it can be realized that the enantioselectivity is maximum when La is used as a lanthanide metal and the chiral chelating agent 1 is used as a ligand (entry 3).

[0016] To further determine the feasibility of the catalytic system, a variety of α,β -unsaturated carbonyl compounds with suitable aldehydes are studied under optimized conditions. A variety of acrylates with diverse steric, geometric, and electronic properties are surveyed in this transformation. These include methyl acrylate (A), t-butyl acrylate (B), phenyl acrylate (C), benzyl acrylate (D), and α -naphthyl acrylate (E). The reaction formula is as follows:



A: $R^2 = CH_3$; B: $R^2 = C(CH_3)_3$; C: $R^2 = Ph$; D: $R^2 = Bn$; and E: $R^2 = \alpha$ -Naphthyl

[0017] The detail steps of the reaction of benzaldehyde and methyl acrylate are described as follows: first, the chiral cheating agent 1 (0.10 g, 0.26 mmol) is dissolved in CH_3CN (2.6 mL), and then $La(OTf)_3$ is added at room temperature. The solution is stirred for 10 min and methyl acrylate (0.22 g, 2.60 mmol), benzaldehyde (0.27 g, 2.60 mmol) and DABCO (0.10 g, 0.78 mmol) are added sequentially. The resulting mixture is stirred for 20 min and the reaction is then quenched with H_2O (5 mL). The resulting solution is extracted with CH_2Cl_2 (10 mL) and the layers are separated. The organic layer is washed with brine, dried with $MgSO_4$ and concentrated. The crude product is purified by silica gel using hexane/EtOAc as the eluent (8/1) to give 0.42 g (88%) of the Baylis-Hillman adduct with 84%ee. The product is analyzed and the data is as follows: $[\alpha]_D = +93.3^\circ$ (c 1.0, $CHCl_3$); lit. $[\alpha]_D = -111.1^\circ$ (c 1.11, MeOH) , for *R* enantiomer; 1H NMR ($CDCl_3$, 200 MHz) δ 7.38-7.26 (m, 5H), 6.32 (d, 1H, $J = 1.0$ Hz), 5.85 (d, 1H, $J = 1.0$ Hz), 5.40 (s, 1H), 3.69 (t, 3H), 3.13 (bs, 1H); ^{13}C NMR ($CDCl_3$, 50 MHz) δ 166.71, 142.06, 141.32, 128.33, 127.72, 126.57, 125.85, 72.89, 51.72; HRMS m/z 192.0777 (calcd for $C_{11}H_{12}O_3$ 192.0786); and HPLC condition: 2-propanol : hexane = 20 : 80 (0.5 mL/min), $t_R = 13.3$ min (*S*) and 15.3 min (*R*).

[0018] The detailed steps of the reaction of benzaldehyde and α -naphthyl acrylate are described as follows: first, a solution of chiral ligand 1 (12 mg, 0.03 mmol) in CH_3CN (2.6 mL) is added $La(OTf)_3$ (9 mg, 0.015 mmol) at room temperature under N_2 atmosphere. The solution is stirred for 10 min and benzaldehyde (54 mg, 0.51 mmol), α -naphthyl acrylate (0.10 g, 0.51 mmol) and DABCO (17 mg, 0.15 mmol) are added sequentially. The resulting mixture is stirred for 20 min and quenched with H_2O (5 mL). The resulting solution is extracted with CH_2Cl_2 (10 mL) and the layers are

separated. The organic layer is washed with brine (10 mL), dried (MgSO_4) and concentrated. The crude product is purified by silica gel using hexane/EtOAc as eluent (8/1) to give 0.13 g (88%) of product as a white solid. The enantiomeric ratios were determined by HPLC analyses using a chiral column. The result is shown in Table 2 as

5 follows:

Table 2. Reaction of acrylates A-E with aldehydes catalyzed by $\text{La}(\text{OTf})_3$ complexes^a

entry	acrylate	R_1CHO ($\text{R}_1 =$)	reaction time/hour	yield(%) ^b	configuration	(%ee) ^c
1	A	CH_3	10	85	<i>S</i>	10
2	A	CH_3CH_2	10	89	<i>S</i>	7
3	A	$(\text{CH}_3)_2\text{CH}$	10	75	<i>S</i>	6
4	A	4-MeOC ₆ H ₄	10	55	<i>S</i>	66
5	B	Ph	10	25	<i>S</i>	70
6	C	Ph	10	97	<i>S</i> ^d	75
7	D	CH_3	10	85	<i>S</i>	65
8	D	CH_3CH_2	10	85	<i>S</i>	65
9	D	Ph	10	75	<i>S</i> ^d	75
10	D	4-MeOC ₆ H ₄	10	50	<i>S</i>	95
11	D	4-NO ₂ C ₆ H ₄	10	93	<i>S</i>	85
12	E	<i>c</i> -C ₆ H ₁₁	1/3	71	<i>S</i>	71 ^e
13	E	CH_3CH_2	1/3	75	<i>S</i>	70 ^f
14	E	Ph	1/3	88	<i>S</i> ^d	81
15	E	4-MeOC ₆ H ₄	1/3	35	<i>S</i>	95
16	E	4-NO ₂ C ₆ H ₄	1/3	82	<i>S</i>	93
17	E	$\text{Ph}(\text{CH}_2)_2\text{CH}_2$	1/3	78	<i>S</i>	81

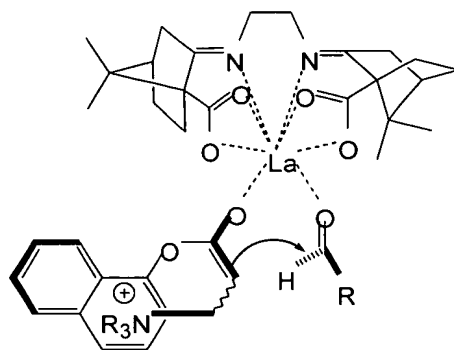
^a All reactions are carried out at room temperature. ^b Isolated yield. ^c Determined by HPLC analysis using a chiral column. ^d The absolute stereochemistry is determined by HPLC analysis after conversion to known product 5A ($R^1 = \text{Ph}$, $R^2 = \text{CH}_3$). The absolute stereochemistry of other products are assigned by analogy, ^e ^f %Ee is
5 determined by conversion to the corresponding ester 5A ($R^1 = \text{c-C}_6\text{H}_{11}$, $R^2 = \text{CH}_3$) for HPLC analysis.

[0019] >From Table 2, it can be realized that the levels of asymmetric induction diminish appreciably when aliphatic aldehydes are treated with methylacrylate under the catalytic conditions employed (Table 2, entries 1-3). Good selectivity is obtained with
10 low chemical yield when t-butyl acrylate was used with benzaldehyde (entry 5). Excellent chemical yield with moderate stereoselectivity was obtained when phenyl acrylate (C) was treated with benzaldehyde (entry 6).

[0020] Moreover, from Table 2, it can be realized that the stereoselectivity is remarkably enhanced when benzyl acrylate (D) was reacted with various aldehydes. A
15 significant improvement in stereoselectivity was obtained when aliphatic aldehydes are reacted with benzyl acrylate (compare entries 1, 2 and 7, 8). The use of aromatic aldehydes with either electron donating or deficient substituents on the benzene ring gave high stereoselectivity (entries 10, 11). The use of benzylacrylate to provide good to high levels of stereoselectivity in the Baylis-Hillman products is interesting. However,
20 the reaction requires 10 hours at room temperature for completion.

[0021] In addition, referring to entries 12 to 17 of Table 2, when α -naphthyl acrylate (E) serves as α,β -unsaturated carbonyl compound, the reaction time is reduced substantially. This represents that the reaction rate is remarkably enhanced. For example, treatment of the cyclohexanecarboxaldehyde with α -naphthyl acrylate

afforded the desired product in 71% yield in 20 min (entry 12). The enantioselectivity was determined to be 71% ee. In the case of aromatic aldehydes, high levels of stereoselectivities were obtained under the same reaction conditions (entries 14-16). The addition of ammonium enolate to an aldehyde is thought to be the rate-determining
 5 step of the Baylis-Hillman reaction. The remarkable rate acceleration using ammonium enolate can be attributed to the potential π -charge stabilization of the intermediates between the naphthalene aromatic system with the azaenolate. Stabilization of the ammonium enolate enhances the addition of DABCO to the α -naphthyl acrylate and therefore accelerates the reaction rates. The proposed mechanism of the
 10 enantioselective Baylis-Hillman reaction is illustrated as follows:



[0022] In conclusion, using the chiral catalysts of this invention catalyzing an asymmetric Baylis-Hillman reaction, a high level of enantioselectivities can be obtained. Maximun enantioselectivities can be obtained using 3 mol% of catalyst. In addition,
 15 when α -naphthyl acrylate is used as a Michael acceptor, the reaction is complete within 20 min, in reasonable chemical yields and enantiomeric excess.

[0023] It would be understood to those skilled in the art that applications of the chiral catalyst of the present invention are not limited to the asymmetric Baylis-Hillman
 20 reaction. The chiral catalyst of the present invention can be applied in other chemical

reactions, for example, a chiral alkylation, a chiral reduction, a chiral cyclization including a [2+2], a [3+2], a [4+2] or a [2+2+2] cyclization, a chiral hydrogenation, a chiral epoxidation, a chiral cyclization of propane, a chiral aziridination, a alkylation, a chiral dialkylation, a chiral hydroxyamination, a chiral amination, an Aldol reaction or a
5 Michael addition reaction.

[0024] It will be apparent to those skilled in the art that various modifications and variations can be made to the structure of the present invention without departing from the scope or spirit of the invention. In view of the foregoing, it is intended that the present invention covers modifications and variations of this invention provided they
10 fall within the scope of the following claims and their equivalents.